Glossary

BMD₁₀ is a Benchmark Dose associated with a 10% response adjusted for background.

Benchmark Response (BMR) is a designated level or percent of response relative to the control level of response used in calculating a BMD

Common Mechanism of Toxicity pertains to two or more pesticide chemicals or other substances that cause a common toxic effect(s) by the same, or essentially the same, sequence of major biochemical events (i.e., interpreted as mode of action).

Comparative effect level (CEL) is a dose by which potency of chemicals may be compared; e.g. the dose causing a maximum of 15% cholinesterase inhibition.

Cumulative Assessment Group (CAG) is a subset of chemicals selected from a common mechanism group for inclusion in a refined quantitative estimate of risk.

Cumulative risk is the risk of a common toxic effect associated with concurrent exposure by all relevant pathways and routes of exposure to a group of chemicals that share a common mechanism of toxicity.

Dose additivity is the Agency's assumption when evaluating the joint risk of chemicals that are toxicologically similar and act at the same target site. In other words, it is assumed that each chemical behaves as a concentration or dilution of every other chemical in the CAG (or chemical mixture). The response of the combination is the response expected from the equivalent dose of an index chemical. The equivalent dose is the sum of the component doses, scaled by each chemical's toxic potency relative to the index chemical.

Index chemical is a chemical used as the point of reference for standardizing the common toxicity of the chemical members of the CAG.

Lowest-Observed-Adverse-Effect Level (LOAEL) is the lowest dose in a toxicity study resulting in adverse health effects

No-Observed-Adverse-Effect Level (NOAEL) is the highest dose in a toxicity study which does not result in adverse health effects

OPCumRisk is a computer program developed at ORD's NHEERL to determine relative potency estimates and PoDs for the index chemical.

Pathway of Exposure is the physical course a pesticide takes from the source to the organism exposed (e.g., through food or drinking water consumption or residential pesticide uses).

Point of Departure (PoD) is a dose that can be considered to be in the range of observed responses, without significant extrapolation. A PoD can be a data point or an estimated point that is derived from observed dose-response data. A PoD is used to mark the beginning of extrapolation to determine risk associated with lower environmentally relevant human exposures.

Relative Potency Factor (RPF) is the ratio of the toxic potency of a given chemical to that of an index chemical in the CAG. Relative potency factors are used to convert exposures of all chemicals in the CAG into their exposure equivalents of the index chemical.

Route of Exposure is the way a chemical enters an organism after contact (e.g., ingestion, inhalation, or dermal absorption).

Steady state inhibition is the time point at which continued dosing at the same level results in no further increase in cholinesterase inhibition.

Acronyms

A Estimate of A (background cholinesterase activity)

AChE Acetylcholinesterase

B Estimate of B (horizontal-asympote from July 2001 analysis)

B/A Ratio of estimate of B/estimate of A

BMD₁₀ A Benchmark Dose associated with a 10% response adjusted for

background

BMDL Lower 95% confidence limit on the BMD₁₀

BMR Benchmark Response -a designated level or percent of response relative

to the control level of response used in calculating a BMD

CEL Comparative effect level - Dose level used to compare potencies

ChEI Cholinesterase inhibition

CL Confidence limit

CNS Central nervous system

D Displacement parameter in expanded model
DER Data evaluation record, a review of a toxicity study

F Female

FIFRA Federal Insecticide, Fungicide, and Rodenticide Act

FQPA Food Quality Protection Act
GOF Model goodness-of-fit
UED Health Effects Division

HED Health Effects Division idose Scaled internal dose

m Estimate of absolute potency for a single cholinesterase measurement in

the July 2001 analysis

IA Log of background cholinesterase activity

*l*m Log slope-scale factor

M Male

MOE Margin of exposure

MRID # Study identification number

NA Not available

NERL National Exposure Research Laboratory

NHEERL National Health and Environmental Effects Laboratory

nlme Non-linear mixed effects model

NOAEL No-Observed-Adverse-Effect Level - the highest dose in a toxicity study

which does not result in adverse health effects

OP Organophosphorous pesticide

OPCumRisk Computer program developed at ORD's NHEERL to determine relative

potency estimates and PoDs for the index chemical.

OPP Office of Pesticide Programs

OPPTS Office of Prevention, Pesticides, and Toxic Substances

ORD Office of Research and Development

P_B Limiting value of minimum cholinesterase activity (horizontal asymptote)

 $\begin{array}{ll} P_{\text{BF}} & \text{Female specific value of } P_{\text{B}} \\ P_{\text{RM}} & \text{Male specific value of } P_{\text{B}} \end{array}$

PBPK Physiologically Based Pharmacokinetics

POD Point of Departure

PNS Peripheral nervous system

RBC Red blood cells

Reference Dose - A dose not expected to cause adverse health effects in RfD

humans

RPF Relative Potency Factor

Shape S

SAP

Scientific Advisory Panel
Transformed horizontal asymptote *t*B